

-continued

Arg	Arg	Pro	Ile	Arg	Asn	Leu	Thr	Phe	Gln	Asp	Leu
1				5					10		

<210> SEQ ID NO 273
 <211> LENGTH: 14
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: VSTB112 minor epitope 1

<400> SEQUENCE: 273

Thr	Trp	Tyr	Arg	Ser	Ser	Arg	Gly	Glu	Val	Gln	Thr	Cys	Ser
1				5				10					

<210> SEQ ID NO 274
 <211> LENGTH: 17
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: VSTB112 minor epitope 2

<400> SEQUENCE: 274

Glu	Ile	Arg	His	His	His	Ser	Glu	His	Arg	Val	His	Gly	Ala	Met	Glu
1				5				10					15		

Leu

<210> SEQ ID NO 275
 <211> LENGTH: 32
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: IGN175A epitope

<400> SEQUENCE: 275

Phe	Lys	Val	Ala	Thr	Pro	Tyr	Ser	Leu	Tyr	Val	Cys	Pro	Glu	Gly	Gln
1				5				10					15		

Asn	Val	Thr	Leu	Thr	Cys	Arg	Leu	Leu	Gly	Pro	Val	Asp	Lys	Gly	His
			20				25					30			

1.-53. (canceled)

54. An antigen-binding molecule which binds to VISTA and inhibits VISTA-mediated signalling, comprising:

(i) a heavy chain variable (VH) region incorporating the following CDRs:

HC-CDR1 having the amino acid sequence of SEQ ID NO:33

HC-CDR2 having the amino acid sequence of SEQ ID NO:34

HC-CDR3 having the amino acid sequence of SEQ ID NO:35;

or a variant thereof, in which one amino acid of HC-CDR1, two amino acids of HC-CDR2 and one amino acid of HC-CDR3 are substituted with another amino acid; and

(ii) a light chain variable (VL) region incorporating the following CDRs:

LC-CDR1 having the amino acid sequence of SEQ ID NO:41

LC-CDR2 having the amino acid sequence of SEQ ID NO:42

LC-CDR3 having the amino acid sequence of SEQ ID NO:43;

or a variant thereof, in which one amino acid of LC-CDR2 is substituted with another amino acid.

55. The antigen-binding molecule according to claim **54**, wherein the antigen-binding molecule comprises:

a VH region comprising an amino acid sequence having at least 70% sequence identity to the amino acid sequence of SEQ ID NO:32; and

a VL region comprising an amino acid sequence having at least 70% sequence identity to the amino acid sequence of SEQ ID NO:40.

56. The antigen-binding molecule according to claim **54**, wherein the antigen-binding molecule is capable of inhibiting interaction between VISTA and a binding partner for VISTA.

57. The antigen-binding molecule according to claim **54**, wherein the antigen-binding molecule is capable of increasing proliferation and/or cytokine production by effector immune cells.

58. A method of treating or preventing a cancer in a subject, the method comprising administering to a subject a therapeutically or prophylactically effective amount of an